California Medical Device Recall Information

Recall Name
Synthes USA Recalls Hemostatic Bone Putty Due to a Potential to Ignite

Recall Date	Product Description	Recalling Firm	Recall Reason
7/05/12	Hemostatic Bone Putty	**Synthes USA, Inc.** West Chester, PA	*Potential to ignite*
Recall Class	**Product Identification**	**Distribution**	**Affected Dates**
I	Hemostatic Bone Putty Part Numbers: • 08.901.001.97S • 08.901.001.98S • 08.901.001.99S • 08.901.001D • VB1025.1 All Lots Recalled	**CA**, nationwide	Manufactured from July 6, 2011 to December 14, 2011

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm316457.htm